K061959

AUG 22 2006

"510(k) Summary"

Submitter's Name:	***MYTECH TECHNOLOGY CO., LTD.***
Address:	*5F.6, Alley 2, Lane 222, Lien Cheng Road,* *Chung Ho City, Taipei Hsien, 235, Taiwan, ROC*
Telephone:	886-2-2247 4816
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Contact Person:	Dr. Jen, Ke-Min
Date Summary Prepared:	7/3/2006
Proprietary Name:	*MYTECH / HAPPY LIFE / MAXIAIDS ARM BLOOD PRESSURE MONITOR, HPL-301, HPL-302, HPL-310, 180310*
Common Name:	BLOOD PRESSURE MONITOR
Classification Name:	NON-INVASIVE BLOOD-PRESSURE MEASUREMENT SYSTEM (per 21CFR section 870.1130)
Device Class:	Class II (performance standards)
Specialty:	CARDIOVASCULAR
Product code:	DXN
Legally Marketed (Predicate) Device :	*MYTECH / HAPPY LIFE ARM BLOOD PRESSURE MONITOR, HPL300 / HPL300A* 510(k) No: **K030221**

Description of the new device:

Basically, the subject device and the predicated device are the same in intended use, technological characteristics, power supply, LCD display, range, accuracy, and operating environments, dimensions, weight, and storage environments. The minor differences are the memory and display layout. That means the predicate device has 60 memory capacities and the new devices have 60 memory capacities. Besides, the new devices add 3 colors LED warming lights for the visional appearance.

The main differences of the new devices are:

- HPL-301 is the representative model of the new devices;
- HPL-302 is same as the HPL-301, add USB port for PC-LINK function;
- HPL-310 is same as the HPL-301, add voice prompt function for arm;
- 180310 is all the same as HPL-310, just using another model name.

Since we also did the relevant electric safety and EMC testing for the new devices, the safety and effectiveness aspects are not raised.

They are decided to be substantially equivalent.

Technological Characteristics of our new device compared to the predicate device:

The technological characteristic of MYTECH / HAPPY LIFE / MAXIAIDS ARM BLOOD PRESSURE MONITOR, HPL-301, HPL-302, HPL-310, 180310 are substantially equivalent to the predicate device. The new devices are of generally the same form and intended to be used in the same manner as the substantially equivalent product; and are passed the relevant EMC and Safety standards. Thus there are substantially equivalent.

Test Summary:

1. **ELECTRIC SAFETY and EMC test reports,**

General safety	*EN 60601-1:1990+A1+A2+A11+A12+A13*	***PASS***
EMC conformity	*EN 60601-1-2: 2001*	***PASS***

2. **WOVEN COTTON SHEETING**

   Uses the 510K Blood-Pressure Cuff

3. **PERFORMANCE & CLINICAL TEST**

   AAMI / ANSI SP10

   The new devices use the same software as the predicate device.

MYTECH TECHNOLOGY CO., LTD. *believes this information and referred document to be sufficient for the FDA to find our proposed device substantially equivalent to the predicate product and other products currently in distribution.*

Jen, Ke-Min

Submitter, 7/3/2006

Official Correspondent

MYTECH TECHNOLOGY CO., LTD.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

AUG 2 2 2006

MYTECH Technology Co., LTD
c/o Dr. Ke-Min Jen
ROC Chinese European Industrial Research Society
No. 58, Fu-Chiun St.
Hsin-Chu City
CHINA (TAIWAN)

Re: K061959
Trade Name: MYTECH HAPPY LIFE/MAXIAIDS Arm Blood Pressure Monitor, Model HPL-301, HPL-302, HPL-310, and 180310
Regulation Number: 21 CFR 870.1130
Regulation Name: Non-Invasive Blood Pressure Measurement System
Regulatory Class: Class II (two)
Product Code: DXN
Dated: July 3, 2006
Received: July 11, 2006

Dear Dr. Ke-Min Jen:

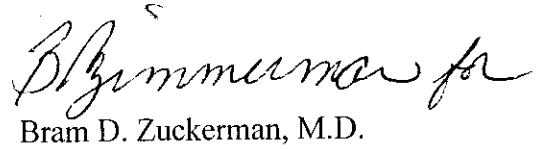
We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.
This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150 or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,



Bram D. Zuckerman, M.D.
Director
Division of Cardiovascular Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

Indications for Use

510(k) Number: K061959

Device Name: ***MYTECH TECHNOLOGY CO., LTD.***

MYTECH / HAPPY LIFE / MAXIAIDS ARM BLOOD PRESSURE MONITOR, HPL-301, HPL-302, HPL-310, 180310

- *Indications for use:*

The MYTECH / HAPPY LIFE / MAXIAIDS Arm Blood Pressure Monitor,, Model HPL-301, HPL-302, HPL-310, 180310 are noninvasive blood pressure measurement systems intended to measure the systolic and diastolic blood pressures and pulse rate of an adult individual, over age 18, at home by using a non-invasive technique in which an inflatable cuff is wrapped around the arm.
The cuff circumference is limited to be 7"-15".

Prescription Use _____ AND/OR Over-The-Counter Use √

(Part 21 CFR 801 Subpart D) (21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Cardiovascular Devices
510(k) Number K061959

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